Appl. No.

: 09/539,032

Filed

March 30, 2000

AMENDMENTS TO THE CLAIMS

1. (Currently amended) A computer-based method for identifying conserved peptide motifs useful as drug targets for use in a host organism, wherein said method comprises the steps of:

- i) computationally generating overlapping peptide sequences from selected <u>pathogenic</u> organisms of length 'N',
- ii) computationally sorting the peptide sequences of length 'N' according to amino acid sequence,
- iii) computationally matching the sorted peptide sequences of length 'N' of the selected <u>pathogenic</u> organisms to produce matched common peptide sequences,
- iv) computationally locating the matched common peptide sequences in the their corresponding protein sequences of step i) to provide locations of said matched common peptide sequences and subsequently labeling the matched common peptide sequences with their origin and location;
 - v) computationally joining overlapping common peptide sequences to obtain extended conserved peptide sequences; and
- vi) comparing said extended conserved peptide sequences obtained in step (v) to host organism protein sequences to determine which of said conserved peptide sequences <u>from said</u> <u>selected pathogenic organisms</u> are not present in host proteins, wherein said conserved peptide sequences which are not present in host proteins are useful as drug targets.
 - 2. (Previously presented) The method of claim 1, wherein 'N' is at least 4.
- 3. (Currently amended) The method of claim 1 wherein the selected <u>pathogenic</u> organisms include at least one of: Mycoplasma pneumoniae, Helicobacter pylori, Hemophilus influenzae, Mycobacterium tuberculosis, Mycoplasma genitalium, Bacillus subtilis, and Escherichia coli.
- 4. (Previously presented) The method of claim 1, wherein the extended conserved peptide sequences comprise one or more of the following sequences:

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1.	AAQSIGEPGTQLT (SEQ ID NO:1)	35.	KMSKSKGN (SEQ ID NO:35)
2.	AGDGTTTAT (SEQ ID NO:2)	36.	KMSKSLGN (SEQ ID NO:36)
3.	AGRHGNKG (SEQ ID NO:3)	37.	KNMITGAAQMDGAIL (SEQ ID NO:37)
4.	AHIDAGKTTT (SEQ ID NO:4)	38.	KPNSALRK (SEQ ID NO:38)
5.	CPIETPEG (SEQ ID NO:5)	39.	LFGGAGVGKTV (SEQ ID NO:39)
6.	DEPSIGLH (SEQ ID NO:6)	40.	LGPSGCGK (SEQ ID NO:40)
7.	DEPTSALD (SEQ ID NO:7)	41.	LHAGGKFD (SEQ ID NO:41)
8.	DEPTTALDVT (SEQ ID NO:8)	42.	LIDEARTPLIISG (SEQ ID NO:42)
9.	DHAGIATQ (SEQ ID NO:9)	43.	LLNRAPTLH (SEQ ID NO:43)
10.	DHPHGGGEG (SEQ ID NO:10)	44.	LPDKAIDLIDE (SEQ ID NO:44)
11.	DLGGGTFD (SEQ ID NO:11)	45.	LPGKŁADC (SEQ ID NO:45)
12.	DVLDTWFSS (SEQ ID NO:12)	46.	LSGGQQQR (SEQ ID NO:46)
13.	ERERGITI (SEQ ID NO:13)	47.	MGHVDHGKT (SEQ ID NO:47)
14.	ERGITITSAAT (SEQ ID NO:14)	48.	NADFDGDQMAVH (SEQ ID NO:48)
15.	ESRRIDNQLRGR (SEQ ID NO:15)	49,	NGAGKSTL (SEQ ID NO:49)
16.	FSGGQRQR (SEQ ID NO:16)	50.	NLLGKRVD (SEQ ID NO:50)
17.	GEPGVGKTA (SEQ ID NO:17)	51.	NTDAEGRL (SEQ ID NO:51)
18.	GFDYLRDN (SEQ ID NO:18)	52.	PSAVGYQPTLA (SEQ ID NO:52)
19.	GHNLQEHS (SEQ ID NO:19)	53.	QRVALARA (SEQ ID NO:53)
20.	GIDLGTTNS (SEQ ID NO:20)	54.	QRYKGLGEM (SEQ ID NO:54)
21.	GINLLREGLD (SEQ ID NO:21)	55.	RDGLKPVHRR (SEQ ID NO:55)
22,	GIVGLPNVGKS (SEQ ID NO:22)	56.	SALDVSIQA (SEQ ID NO:56)
23.	GKSSLLNA (SEQ ID NO:23)	57.	SGGLHGVG (SEQ ID NO:57)
24.	GLTGRKIIVDTYG (SEQ ID NO:24)	58.	SGSGKSSL (SEQ ID NO:58)
25.	GPPGTGKTLLA (SEQ ID NO:25)	59.	SGSGKSTL (SEQ ID NO:59)
26.	GPPGVGKT (SEQ ID NO:26)	60.	SVFAGVGERTREGND (SEQ ID NO:60)
27.	GSGKTTLL (SEQ ID NO:27)	61.	TGRTHQIRVH (SEQ ID NO:61)
28.	GTRIFGPV (SEQ ID NO: 28)	62.	TGVSGSGKS (SEQ ID NO:62)
29.	IDTPGHVDFT (SEQ ID NO:29)	63.	TLSGGEAQRI (SEQ ID NO: 63)
30.	ILAHIDHGKSTL (SEQ ID NO:30)	64.	TNKYAEGYP (SEQ ID NO:64)
31.	INGFGRIGR (SEQ ID NO:31)	65.	TPRSNPATY (SEQ ID NO:65)
32.	IREGGRTVG (SEQ ID NO:32)	66.	VEGDSAGG (SEQ ID NO:66)
33.	IVGESGSGKS (SEQ ID NO:33)	67.	VRKRPGMYIG (SEQ ID NO:67)
34.	KFSTYATWWI (SEQ ID NO:34)		

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5. (Canceled)

6. (Previously presented) The method of any one of claims 1-4 wherein the conserved peptide sequences are found within the sequences of at least one of the following proteins:

I DNA DIRECTED RNA POLYMERASE BETA CHAIN

II EXONUCLEASE ABC SUBUNIT A

III EXONUCLEASE ABC SUBUNIT B

IV DNA GYRASE SUBUNIT B

V ATP SYNTHASE BETA CHAIN

VI S-ADENOSYLMETHIONINE SYNTHETASE

VII GLYCERALDEHYDE 3-PHOSPHATE DEHYDROGENASE

VIII ELONGATION FACTOR G (EF-G)

IX ELONGATION FACTOR TU (EF-TU)

X 30S RIBOSOMAL PROTEIN S12

XI 50S RIBOSOMAL PROTEIN L12

XII 50S RIBOSOMAL PROTEIN L14

XIII VALYL tRNA SYNTHETASE

XIV CELL DIVISION PROTEIN FtSH HOMOLOG

XV DnaK PROTEIN (HSP70)

XVI GTP BINDING PROTEIN LepA

XVII OLIGOPEPTIDE TRANSPORT ATP BINDING PROTEIN OPPF.

7. (Previously presented) The method of claim 1, wherein step (iii) comprises: selecting organism names from a menu;

iteratively comparing peptide sequences of a first organism to peptide sequences of a second organism; and, for matching sequences, writing sequences to a first file for the first organism and to a second file for the second organism.

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8. (Currently amended) The method of claim 1 wherein step (iv) comprises: selecting protein sequences; iteratively comparing matched peptide sequences to protein sequences; and if the peptide is found in a protein sequence, labeling the peptide sequence in a file associated with the protein with: a) a protein identification number (PID), b) a location in the protein sequence, and c) a name of the selected a pathogenic organism chosen from the group of selected pathogenic organisms of step iii).

9. (Previously presented) The method of claim 1, wherein step (v) comprises: iteratively comparing peptide sequences on matched peptide locations; determining overlapping matched peptides; and determining extended peptide sequences based on overlapping matched peptide sequences.

10-12. (Canceled)